A single copy subclone, p1-101, from cosmid 3-3B, defines three RFLPs on 10pter-q23 [HGM9 no. D10S4]

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SOURCE AND DESCRIPTION OF CLONE: p1-101, a 480 bp Sau3A fragment subcloned into pJB8 from cosmid 3-3B, isolated from a random human genomic library.

POLYMORPHISMS: TaqI identifies a 2-allele RFLP with A1= 7.4, A2=4.9 kb. SacI identifies two 2-allele RFLPs with B1=4.7, B2=0.81 kb and C1=3.2, C2=1.75 kb.

ALLELE FREQUENCIES: studied at least 46 European Caucasians:

A1=0.56±.051 B1=0.39±.043 C1=0.81±.035 A2=0.44±.051 B2=0.61±.043 C2=0.19±.035

NOT POLYMORPHIC FOR: Mspl, EcoRI, HindIII, BamHI, Psti, PvuII and Rsal

CHROMOSOMAL LOCALISATION: 1 Opter-q23, by use of somatic cell hybrid panels.

MENDELIAN INHERITANCE: At locus A, codominant inheritance has been shown in 20 informative families with a total of 63 children. At loci B and C, codominant inheritance has been shown in 3 informative families with a total of 23 children.

PROBE AVAILABILITY: Available for collaboration.

OTHER COMMENTS: Twenty-seven of 31 unrelated individuals (87%) typed with both Taql and Sacl were heterozygous at one or more of the three loci.

REFERENCE: Litt, M. and Litt, R (1986) Am J Hum Genet 39:A161.

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